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**THE UNITED STATES DISTRICT COURT**  
**DISTRICT OF UTAH**

Abby Jergins, an individual; Bill Jergins,  
an individual; and Archi Hipkins, an  
individual; on behalf of themselves and all  
others similarly situated,

Plaintiffs,

vs.

Johnson & Johnson Consumer, Inc., a  
New Jersey corporation; The Procter &  
Gamble Company, an Ohio corporation;  
GlaxoSmithKline Consumer Healthcare  
Holdings (US) LLC, a Delaware limited  
liability company; and Haleon US Capital  
LLC, a Delaware limited liability  
company;

and

John Does 1-200,

Defendants.

Case No.

**CLASS ACTION COMPLAINT**

**JURY DEMANDED**

**COMPLAINT**

Plaintiffs Abby Jergins, Bill Jergins, and Archi Hipkins (collectively “Plaintiffs”), and by and through their undersigned counsel, hereby bring this action on behalf of themselves and all others similarly situated, against Defendants, Johnson & Johnson Consumer, Inc.; The

Procter & Gamble Company; GlaxoSmithKline Consumer Healthcare Holdings (US) LLC; Haleon US Capital LLC; and Does 1 through 200 (collectively, “Defendants”), and state;

### **INTRODUCTION**

1. This is a class action for damages related to Defendants’ wrongful conduct in connection with the marketing, advertising, promoting, distribution and sale of products containing phenylephrine—a purported decongestant used as an active ingredient in at least 250 products, including without limitation Sudafed, Vicks Dayquil & Nyquil, Tylenol Cold & Flu, Advil Cold & Sinus, and many others, including generic brands developed by major retailers like CVS, Walmart, Target and Walgreens (the “Phenylephrine Products”).

2. Defendants manufacture, test, promote, advertise, market, distribute and sell the Phenylephrine Products for the treatment of congestion and other associated cold and flu symptoms. Millions of Utahns, and hundreds of millions of Americans, spend hard-earned money to purchase these products for help relieving congestion and other associated cold and flu symptoms because they are told by the above-captioned Defendants that they work for that very purpose.

3. For years, Defendants have advertised and marketed the Phenylephrine Products to unsuspecting consumers despite knowing that phenylephrine is ineffective for the treatment of nasal congestion and the other cold and flu symptoms for which Defendants promote its use. On or about September 12, 2023, the Federal Drug Administration, after careful study and consideration, announced publicly that phenylephrine is ineffective as a treatment for such symptoms.

4. As a proximate result of Defendants’ deceptive, fraudulent, unlawful, and/or unfair conduct, Plaintiffs and the putative class collectively suffered hundreds of millions of dollars in damages in reliance upon Defendants’ knowingly false representations about the effectiveness of phenylephrine and the Phenylephrine Products.

5. Plaintiffs therefore demand judgment against Defendants and request, among other things, compensatory damages, statutory damages, punitive damages, attorneys’ fees,

costs and all other available remedies and damages allowed by law.

### **PARTIES**

#### **a. Plaintiffs**

6. At all relevant times, Plaintiff **Abby Jergins** was and has been a resident and citizen of the State of Utah and a resident of the City of St. George.

7. On numerous occasions within the statutory time period, in reliance upon Defendants' intentionally false and fraudulent marketing, Plaintiff Abby Jergins purchased the Phenylephrine Products, specifically including, Sudafed, Vicks Nyquil & Dayquil, and Tylenol Cold & Flu, each of which contained phenylephrine, within the State of Utah for the treatment of cold and flu symptoms.

8. At all relevant times, Plaintiff **Bill Jergins** was and has been a resident and citizen of the State of Utah and a resident of the City of St. George.

9. On numerous occasions within the statutory time period, in reliance upon Defendants' intentionally false and fraudulent marketing, Plaintiff Abby Jergins purchased the Phenylephrine Products, specifically including, Sudafed, Vicks Nyquil & Dayquil, and Tylenol Cold & Flu, each of which contained phenylephrine, within the State of Utah for the treatment of cold and flu symptoms.

10. At all relevant times, Plaintiff **Archi Hipkins** was and has been a resident and citizen of the State of Utah and a resident of the City of Sandy.

11. On numerous occasions within the statutory time period, in reliance upon Defendants' intentionally false and fraudulent marketing, Plaintiff Archi Hipkins purchased the Phenylephrine Products, specifically including, Advil Cold & Sinus, which contained phenylephrine, within the State of Utah for the treatment of cold and flu symptoms.

#### **b. Defendants**

12. Defendant **Johnson & Johnson Consumer, Inc.** is a New Jersey corporation, with headquarters and a principal place of business in the State of New Jersey. Upon information and belief, Defendant Johnson & Johnson Consumer, Inc. is a wholly owned

subsidiary of Johnson & Johnson, a New Jersey corporation, with headquarters and a principal place of business in the State of New Jersey (collectively “J&J”). At all times relevant to this complaint, Defendant J&J was engaged in the business of manufacturing, marketing, testing, promoting, selling, and/or distributing certain of the Phenylephrine Products, including but not limited to, Tylenol, Sudafed, and Benadryl.

13. Defendant **The Procter & Gamble Company** (“Procter”) is an Ohio corporation with headquarters and principal place of business in the State of Ohio. At all times relevant to this complaint, Defendant Procter was engaged in the business of manufacturing, marketing, testing, promoting, selling, and/or distributing certain of the Phenylephrine Products, including but not limited to, Vicks Dayquil & NyQuil.

14. Defendant **GlaxoSmithKline Consumer Healthcare Holdings (US) LLC** is a Delaware limited liability company with its headquarters and principal place of business in the State of New Jersey. Upon information and belief, GlaxoSmithKline Consumer Healthcare Holdings (US) LLC is a wholly-owned subsidiary of GlaxoSmithKline PLC a public limited company organized under the laws of England and Wales (collectively “GSK”). At all times relevant to this complaint, Defendant GSK was engaged in the business of manufacturing, marketing, testing, promoting, selling, and/or distributing certain of the Phenylephrine Products, including but not limited to, Advil Cold & Sinus.

15. Defendant **Haleon US Capital LLC** is a Delaware limited liability company with its headquarters and principal place of business in the State of New Jersey. Upon information and belief, Haleon US Capital LLC is a wholly-owned subsidiary of Haleon PLC a public limited company organized under the laws of England and Wales (collectively “Haleon”). At all times relevant to this complaint, Defendant Haleon was engaged in the business of manufacturing, marketing, testing, promoting, selling, and/or distributing certain of the Phenylephrine Products, including but not limited to, Advil Cold & Sinus.

16. The true names and capacities of defendants Does 1 through 200 are currently unknown to Plaintiffs who, therefore, sue these defendants under these fictitious names. These

defendants are each directly and/or vicariously responsible, in some manner, for the harms alleged herein. If/when Plaintiffs learn these defendants' true names and capacities, Plaintiffs will seek leave to amend this pleading accordingly.

17. The true names and/or capacities, whether individual, corporate, partnership, associate, governmental, or otherwise, of Defendants Does 1 through 200, inclusive, and each of them, are unknown to Plaintiffs at this time, who therefore sues said Defendants by such fictitious names. Plaintiffs are informed and believe, and thereon allege, that each Defendant designated herein as a Doe caused injuries and damages proximately thereby to Plaintiffs as hereinafter allege; and that each Doe defendant is liable to Plaintiffs for the acts and omissions alleged herein below, and the resulting injuries to Plaintiffs, and damages sustained by Plaintiffs. Plaintiffs will amend this Complaint to allege the true names and capacities of said Doe Defendants when that same is ascertained.

#### **JURISDICTION & VENUE**

18. This Court has original jurisdiction pursuant to 28 U.S.C. § 1332(d)(2). The matter in controversy, exclusive of interest and costs, exceeds the sum or value of \$5,000,000 and is a class action in which there are in excess of 100 class members and many members of the class are citizens of a state different from Defendants.

19. This Court has personal jurisdiction over Defendants, which are authorized to conduct and do conduct business in Utah. Defendants have engaged in the business of designing, developing, manufacturing, testing, packaging, promoting, marketing, distributing, and/or selling the Phenylephrine Products to Plaintiffs in Utah and Defendants have sufficient minimum contacts with this State and/or sufficiently avail themselves of the markets in this State through their promotion, sales, distribution and marketing within the State to render exercise of jurisdiction by this Court permissible.

20. Venue is proper in this Court pursuant to 28 U.S.C. §§ 1391(a) and (b) because a substantial part of the events or omissions giving rise to Plaintiffs' claims occurred while they resided in this judicial district. Venue is also proper under 18 U.S.C. § 1965(a) because the

Defendants transact substantial business in this District.

**CLASS ACTION ALLEGATIONS**

21. Pursuant to Rules 23(a), (b)(2), (b)(3), and (c)(4) of the Federal Rules of Civil Procedure, Plaintiffs bring this class action on their own behalf and on behalf of all other similarly situated consumers in the United States as members of the following proposed Nationwide and Utah State classes. The proposed Classes are defined as follows:

- a) **Nationwide Class:** During the fullest period allowed by law, all persons within the United States who purchased the Phenylephrine Products, or any of them, at any time and at any location (the “Class”).
- b) **Utah Subclass:** During the fullest period allowed by law, all persons who, while a resident of Utah, purchased the Phenylephrine Products at any location in Utah, including without limitation any online purchase made from Utah (regardless of the shipping address of the consumer) (the “Utah Subclass” or the “Subclass”).
- c) Nationwide class and Utah Subclass members are collectively referred herein as “Class Members.”
- d) Like Plaintiffs, all Class Members purchased the Phenylephrine Products based on the misrepresentations that said products were effective in the treatment of congestion and other associated cold and flu symptoms, and that such understanding was reasonable and was a material basis for the decision to purchase the Phenylephrine Products, which Defendants intended to foster through its various marketing activities in connection with the sale of the Phenylephrine Products

22. Excluded from the Class and Subclass are assigned judges and members of their families within the first degree of consanguinity, Defendants, and their subsidiaries, affiliates, officers, and directors.

23. Excluded from the Class and Subclass are individuals who allege personal bodily

injury resulting from the use of Phenylephrine Products.

24. The requirements of Federal Rule of Civil Procedure 23 are satisfied for the Class and Utah Subclass.

25. The proposed Class and Utah Subclass are so numerous that individual joinder of all their members is impracticable because members of the Class number in the tens or hundreds of thousands. The precise number of Class members and their identities are unknown to Plaintiffs at this time but are objectively ascertainable and will be determined through appropriate discovery.

26. Defendants possess objective evidence as to the identity of each Class Member and, to a reasonable degree of certainty, the damages suffered by each Class Member, including without limitation sales receipts, phone numbers, names, rewards accounts data, credit card data, customer service complaint forms/emails/date, and other evidence which objectively identifies class members.

27. Class Members may be notified of the pendency of this action by mail, publication and/or through the records of Defendants and third-party retailers and vendors.

28. There are common questions of law and fact affecting Plaintiffs and Class Members. Common legal and factual questions include, but are not limited to:

- a) Whether Defendants market and advertise the Phenylephrine Products in a way that is false or misleading.
- b) Whether by the misconduct set forth in this complaint, Defendants have engaged and continue to engage in unfair, fraudulent, or unlawful business practices;
- c) Whether Defendants' conduct was committed knowingly and/or intentionally;
- d) Whether Defendants' conduct constitutes violations of the federal and/or state laws asserted herein;
- e) Whether Defendants had a duty to correct their fraudulent statements;

- f) Whether Plaintiffs and Class members were harmed by Defendants' false statements;
- g) Whether Defendants were unjustly enriched by their conduct;
- h) Whether Plaintiffs and Class members are entitled to punitive damages;
- i) Whether the Plaintiffs and Class members are entitled to recover statutory attorney's fees;
- j) Whether, as a result of Defendants' misconduct as alleged herein, Plaintiffs and Class Members are entitled to restitution, injunctive and/or monetary relief and, if so, the amount and nature of such relief.

29. Plaintiffs' claims are typical of the claims of the proposed Class and Subclass because Plaintiffs and Class Members were harmed in the same manner by the same conduct.

30. Plaintiffs and Class Members have all sustained economic injury arising out of Defendants violations of common and statutory law alleged herein.

31. Plaintiffs will fairly and adequately represent and protect the interests of the Class and Subclass.

32. Plaintiffs' interests do not conflict with the interests of the Class and Subclass they seek to represent. Plaintiffs have retained counsel competent and experienced in prosecuting class actions, and Plaintiffs intend to prosecute this action vigorously.

33. The class mechanism is superior to other available means for the fair and efficient adjudication of the claims of Plaintiffs and Class Members.

34. Given the relatively small amount of damages at stake for any of the individual Class Members, individual litigation is not practicable.

35. Individual Class Members will not wish to undertake the burden and expense of individual cases.

36. In addition, individualized litigation increases the delay and expense to all parties and multiplied the burden on the judicial system. Individualized ligation also presents the potential for inconsistent or contradictory judgments.



37. In contrast, the class action device presents far fewer management difficulties and provides the benefits of single adjudication, economy of scale, and comprehensive supervision by a single court.

38. Questions of law and fact common to all Class Members predominate over any questions affecting only individual Class Members. Injuries sustained by Plaintiffs and Class Members flow, in each instance, from a common nucleus of operative facts as set forth above.

39. In each case, Defendant used deceptive marketing and sales techniques aimed at the Class Members, causing harm to all Class Members as a result of such intentional conduct. The resolution of these central issues will be the focus of the litigation and predominate over any individual issues.

40. Proposed class counsel possesses the knowledge, experience, reputation, ability, skill, and resources to represent the class and should be appointed lead counsel for the class.

### **TOLLING OF STATUTE OF LIMITATIONS**

#### **a. Discovery Rule Tolling**

41. As a result of the acts and omissions of Defendants, Plaintiffs could not have discovered, through the exercise of reasonable due diligence, that the active ingredient in the Phenylephrine Products was ineffective, as has now been declared by the Federal Drug Administration. Thus, the applicable limitations periods did not begin to accrue until Plaintiffs discovered, or through the exercise of reasonable diligence should have discovered, Defendants' wrongful acts and omissions.

#### **b. Fraudulent Concealment Tolling**

42. All applicable statutes of limitation have also been tolled by Defendants' knowing and active fraudulent concealment and misrepresentations about the effectiveness of phenylephrine and the Phenylephrine Products throughout the time period relevant to this action.

43. Defendants are under a continuing duty to disclose the true character, quality, efficacy, safety issues and safety concerns of phenylephrine and the Phenylephrine Products to

its users, including Plaintiffs specifically. To date, Defendants have nevertheless failed to adequately and fully inform consumers about these matters, as discussed above.

44. Plaintiffs reasonably relied upon Defendants' knowing, affirmative misrepresentations and/or active concealment when Plaintiffs—and millions of similarly-situated Utahns and Americans—purchased the Phenylephrine Products based on the representations and advertisements touting the effectiveness of such products in the treatment of congestion and other associated cold and flu symptoms.

45. Because Defendants actively concealed the true facts about the ineffectiveness of phenylephrine and the Phenylephrine Products, they are estopped from relying on any statutes of limitations defense.

### **CAUSES OF ACTION**

#### **FIRST CAUSE OF ACTION**

##### **Fraudulent Misrepresentation**

46. Plaintiffs reallege and incorporate the allegations made above as if fully set forth below.

47. Plaintiffs bring this claim individually and on behalf of the Class.

48. At all relevant times, Defendants had the duty and obligation to truthfully represent to Plaintiffs the facts concerning the ineffectiveness of phenylephrine and the Phenylephrine Products. Instead, Defendants aggressively (and falsely) advertised the effectiveness of phenylephrine and the Phenylephrine Products, despite the fact that each such Defendant knew that phenylephrine and the Phenylephrine Products were entirely ineffective against congestion and the associated cold & flu symptoms the Phenylephrine Products were advertised to treat.

49. Defendants willfully deceived Plaintiffs and the public in general by making these intentional misrepresentations regarding the efficacy of phenylephrine and the Phenylephrine Products.

50. At the time the aforesaid misrepresentations were made, Defendants intended to

induce Plaintiffs to rely upon such misrepresentations.

51. At the time Defendants made the above-described misrepresentations, Plaintiffs and the public in general, reasonably believed them to be true. In reasonable and justified reliance upon said misrepresentations, Plaintiffs purchased the Phenylephrine Products.

52. As a direct and proximate result of Defendants' conduct, Plaintiffs and the Class suffered serious financial harm, including the expenditure of substantial sums to purchase the Phenylephrine Products, which Defendants knew were and are ineffective for their advertised purpose.

## **SECOND CAUSE OF ACTION**

### **Negligent Misrepresentation**

53. Plaintiffs reallege and incorporate the allegations made above as if fully set forth below.

54. Plaintiffs bring this claim individually and on behalf of the Class.

55. At all relevant times, Defendants had the duty and obligation to truthfully represent to Plaintiffs the facts concerning the ineffectiveness of phenylephrine and the Phenylephrine Products. Instead, Defendants aggressively (and falsely) advertised the effectiveness of phenylephrine and the Phenylephrine Products, despite the fact that each such Defendant should have known that phenylephrine and the Phenylephrine Products were entirely ineffective against congestion and the associated cold & flu symptoms the Phenylephrine Products were advertised to treat.

56. Defendants recklessly or at least negligently deceived Plaintiffs and the public in general by making these misrepresentations regarding the efficacy of phenylephrine and the Phenylephrine Products.

57. At the time the aforesaid misrepresentations were made, Defendants understood that their careless misrepresentations would induce Plaintiffs to rely upon them.

58. At the time Defendants made the above-described misrepresentations, Plaintiffs and the public in general, reasonably believed them to be true. In reasonable and justified

reliance upon said misrepresentations, Plaintiffs purchased the Phenylephrine Products.

59. As a direct and proximate result of Defendants' conduct, Plaintiffs and the Class suffered serious financial harm, including the expenditure of substantial sums to purchase the Phenylephrine Products, which Defendants knew or should have known were and are ineffective for their advertised purpose.

### **THIRD CAUSE OF ACTION**

#### **Breach of Express Warranty**

60. Plaintiffs reallege and incorporate the allegations made above as if fully set forth below.

61. Plaintiffs bring this claim individually and on behalf of the Class.

62. Section 2-313 of the Uniform Commercial Code provides that an affirmation of fact or promise, including a description of the goods, becomes part of the basis of the bargain and creates an express warranty that the goods shall conform to the promise and to the description.

63. At all times, Utah and other states have codified and adopted the Uniform Commercial Code governing the express warranty of merchantability.

64. Plaintiffs and each member of the Class formed a contract with Defendants at the time Plaintiffs and the other members of the Class purchased the Phenylephrine Products. The terms of that contract include the cognitive health benefit promises and affirmations of fact made by Defendants through each of their marketing and advertising of the Phenylephrine Products as described herein. These representations constitute express warranties, became part of the basis of the bargain, and are part of a standardized contract between Plaintiffs and the members of the Class on the one hand, and Defendants on the other.

65. All conditions precedent to Defendants' liability under this contract have been performed by Plaintiffs and the Class Members.

66. At all relevant times, Defendants had the duty and obligation to truthfully represent to Plaintiffs the facts concerning the ineffectiveness of phenylephrine and the

Phenylephrine Products. Instead, Defendants aggressively (and falsely) advertised the effectiveness of phenylephrine and the Phenylephrine Products, despite the fact that each such Defendant knew that phenylephrine and the Phenylephrine Products were entirely ineffective against congestion and the associated cold & flu symptoms the Phenylephrine Products were advertised to treat.

67. Defendants breached the terms of this contract, including the express warranties, with Plaintiffs and the Class by not providing the Phenylephrine Products that could provide the cognitive health benefits as represented and described above.

68. As a result of Defendants' breach of their warranty, Plaintiffs and the Class have been damaged in the amount of the purchase price of the Phenylephrine Products they purchased.

#### **FOURTH CAUSE OF ACTION**

##### **Strict Liability-Design and Manufacturing Defect**

69. Plaintiffs reallege and incorporate the allegations made above as if fully set forth below.

70. Plaintiffs bring this claim individually and on behalf of the Class.

71. At the time that the Phenylephrine Products left the control of the Defendants, the Phenylephrine Products were defective as a result of Defendants' design, manufacture, alteration, or modification. The defects included, but are not limited to, materials that are unsafe for human contact, and/or materials not identified on the Product itself.

72. At all relevant times, Defendant knew and intended that the Phenylephrine Products would be purchased and used by members of the general public who would rely on Defendants to properly identify the relevant characteristics and usefulness of the Product.

73. At the time of the incidents giving rise to this Complaint, the Phenylephrine Products were being used in a manner that was foreseeable by the Defendants and in a manner which the Phenylephrine Products were intended to be used.

74. Defendants knew or should have known their manufacture or design of the

Phenylephrine Products was defective, causing the Phenylephrine Products to fail to perform as an ordinary consumer would expect when used in an intended or reasonably foreseeable manner.

75. In addition, the risks inherent in the design of the Phenylephrine Products outweighs any benefits of that design.

76. As a direct and proximate result of Defendants' conduct, Plaintiffs and the Class have suffered and continue to suffer serious harm.

### **FIFTH CAUSE OF ACTION**

#### **Violation of the Utah Consumer Sales Practices Act (Utah Code Ann. § 13-11-1, *et seq.*)**

#### **(On behalf of Plaintiffs and the Utah Subclass Members)**

77. Plaintiffs reallege and incorporate the allegations made above as if fully set forth below.

78. Plaintiffs assert this Fifth Cause of Action on behalf of themselves and all other similarly-situated persons in Utah that paid hard-earned money for the Phenylephrine Products based on Defendants' deceptive, false, unfair and unlawful marketing strategy touting the effectiveness of phenylephrine and the Phenylephrine Products for treatment of congestion and associated cold and flu symptoms.

79. At all times relevant herein, Defendants were and are each a "person" as defined in Utah Code Ann. § 13-11-3(5), and a "supplier" as defined in Utah Code Ann. § 13-11-3(6).

80. By engaging in the above-described conduct, Defendants, and each of them, acted in a manner that is unlawful, deceptive, unfair, and fraudulent, and have thus engaged in unfair business practices to the extreme detriment of Plaintiffs, which conduct is prohibited under the Utah Consumer Sales Practices Act (Utah Code Ann. §13-11-1, *et seq.*)("UCSPA").

81. Defendants have acted unfairly and deceptively, in violation of the UCSPA, by knowingly and fraudulently advertising to consumers, including Plaintiffs, that phenylephrine and its Phenylephrine Products were effective against congestion and the associated cold & flu symptoms. This representation was likely to mislead consumers acting reasonably under the

circumstances, and did mislead consumers acting reasonably under the circumstances, including Plaintiffs.

82. Defendants' conduct has caused Plaintiffs and the Subclass to suffer harm, including through the payment of monies for the purchase of the Phenylephrine Products.

**Additional Allegations Regarding Punitive Damages**

**(All Applicable Causes of Action)**

83. The acts and omissions of Defendants described herein consisted of oppression, fraud and/or malice and were done with advance knowledge, conscious disregard of the rights of others and/or ratification by Defendants' officers, directors and/or managing agents.

84. Defendants' actions amounted to actual malice or reckless indifference to the likelihood of harm associated with their acts and omissions.

85. Plaintiffs are entitled to punitive damages because Defendants misled, misrepresented and/or withheld information and materials from consumers and the public at large, including Plaintiffs, concerning the efficacy of phenylephrine and the Phenylephrine Products.

86. Despite the fact that Defendants were or should have been in possession of evidence demonstrating the ineffectiveness of phenylephrine and the Phenylephrine Products, Defendants continued to market Phenylephrine Products by providing false and misleading information with regard to the efficacy of such products.

87. Defendants failed to provide consumers, including Plaintiffs, with available materials, information and warnings that would have ultimately dissuaded them from purchasing and consuming such products, thus depriving otherwise uninformed consumers from weighing the true risks and benefits of purchasing and ingesting the Phenylephrine Products.

88. Defendants' conduct was committed with knowing, conscious and deliberate disregard for the rights and safety of consumers, including Plaintiffs, thereby entitling Plaintiffs to punitive damages in an amount appropriate to punish the Defendants and deter them from

similar conduct in the future.

89. Consequently, Defendants are liable for punitive damages in an amount to be determined by the jury at trial.

**PRAYER FOR RELIEF**

WHEREFORE, Plaintiffs, on behalf of themselves and the putative Class Members, prays for a judgment:

- a. Certifying the Class and the Utah Subclass as requested herein, and appointing Plaintiffs and their counsel to represent the Class and the Utah Subclass.
- b. Awarding Plaintiffs and the proposed Class Members damages;
- c. Awarding restitution and disgorgement of Defendants' revenues to Plaintiffs and the proposed Class Members;
- d. Awarding declaratory and injunctive relief as permitted by law or equity, including: enjoining Defendants from continuing the unlawful practices as set forth herein, and directing Defendants to identify, with Court supervision, victims of its conduct and pay them all money it is required to pay;
- e. Ordering Defendants to engage in a corrective advertising campaign;
- f. Awarding punitive damages;
- g. Awarding actual damages in favor of Plaintiffs and all members of the proposed Utah Subclass;
- h. Awarding the costs and expenses of this litigation to Plaintiffs;
- i. Awarding reasonable attorneys' fees and costs to Plaintiffs as provided by law;
- j. Awarding pre-judgment and post-judgment interest to Plaintiffs; and
- k. For such further relief as this Court deems necessary, just and proper.



**DEMAND FOR JURY TRIAL**

Plaintiffs hereby demands a trial by jury on all issues so triable.

Respectfully Submitted,

**SINGLETON SCHREIBER, LLP**

/s/ John Ternieden  
By: JOHN TERNIEDEN, SBN 18740

Attorneys for Plaintiffs